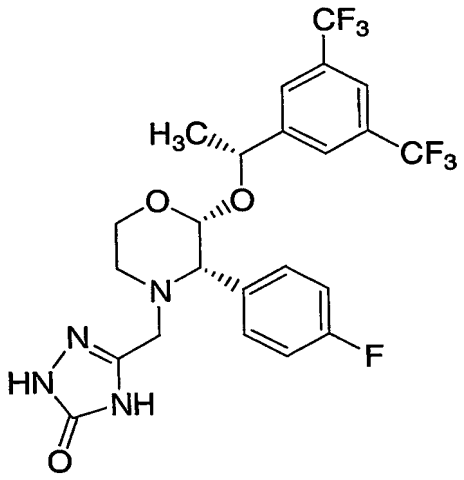
TITLE OF THE INVENTION

PROCESS FOR 5-[[2(R)-[1(R)-[3,5-BIS(TRIFLUOROMETHYL)PHENYL] ETHOXY]-3(S)-(4-FLUOROPHENYL)-4-MORPHOLINYL]METHYL]-1,2-DIHYDRO-3H-1,2,4-TRIAZOL-3-ONE

BACKGROUND OF THE INVENTION

The present invention relates to processes for the preparation of 5-[[2(R)-[1(R)-[3,5-bis(trifluoromethyl)phenyl]ethoxy]-3(S)-(4-fluorophenyl) -4-morpholinyl]methyl]-1,2-dihydro-3H-1,2,4-triazol-3-one, aprepitant,

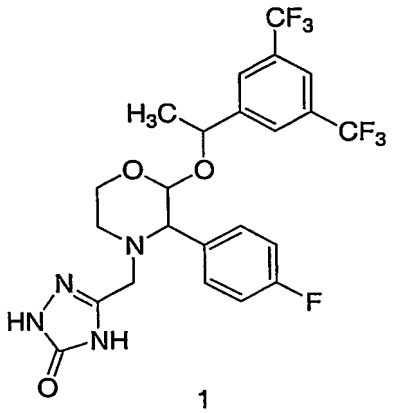


which is a useful therapeutic agent, specifically as a substance P (neurokinin-1) receptor antagonist. This compound is disclosed as having therapeutic utility in U.S. Patent No. 5,719,147.

U.S. Patent Nos. 5,637,699, 6,096,742, 6,229,010 and 6,297,376 relate to processes of manufacture and the discovery of polymorphic forms of this compound. In contrast to previously known processes, the present invention provides a more practical and economical method for preparing the compound in relatively high yield and purity. As such, there is a need for a process for the preparation of the compound that is cost-effective and utilizes readily available reagents.

SUMMARY OF THE INVENTION

The present invention relates to a process for preparing a compound of formula 1:



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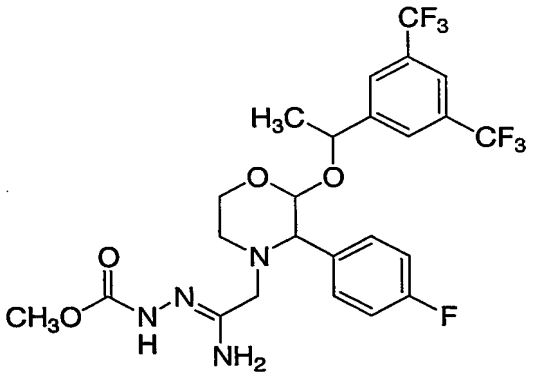
comprising:

cyclizing a compound of formula 4:



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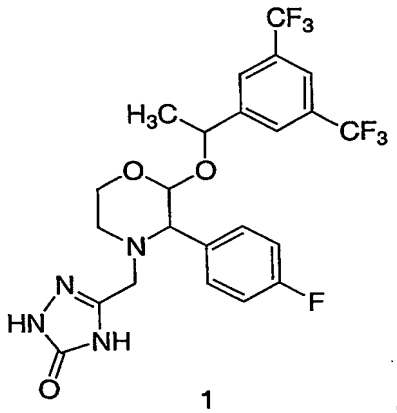
at a temperature of 140-150°C to produce the compound of formula 1.

In particular, such compounds are substance P (neurokinin-1) receptor antagonists which are useful, e.g., in the treatment of psychiatric disorders, inflammatory diseases and emesis.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to a process for preparing a compound of formula 1:



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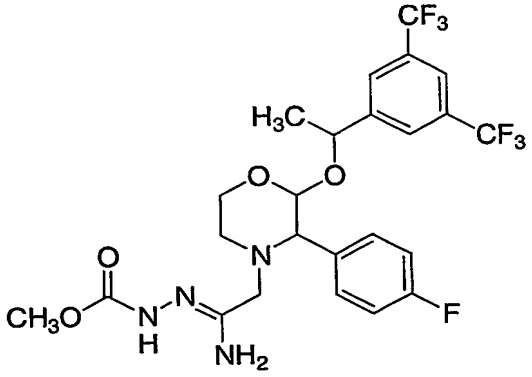
The process comprises :

cyclizing a compound of formula 4:



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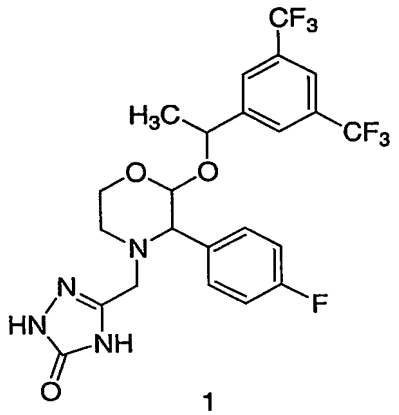
at a temperature of 140-150°C to produce the compound of formula 1.

More particularly, the present invention is directed to processes for the preparation of a compound of formula 1:



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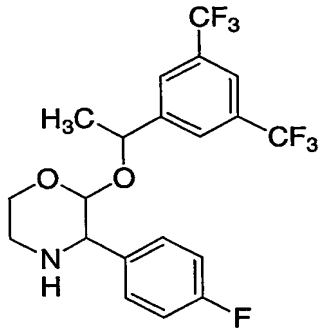
The processes are comprised of:

(a) reacting the hydrochloride salt of a compound of formula 2:



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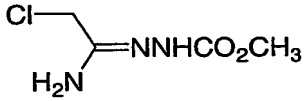
in the presence of an inorganic base and toluene with a compound of the formula 3:



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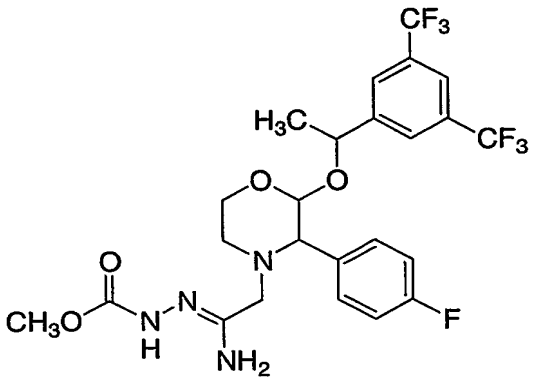
to produce the compound of formula 4:



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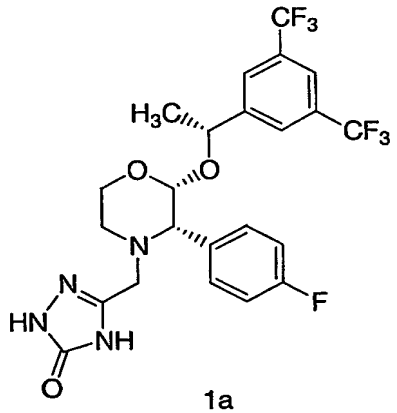
(b) washing with an aqueous phase, and

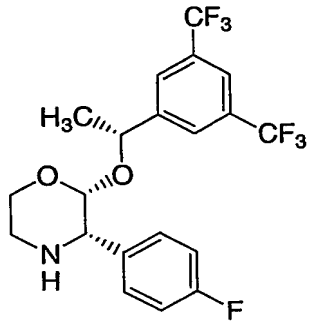
(c) cyclizing at a temperature of 140-150°C to produce the compound of formula 1.

Even more particularly, a process for preparing a compound of formula 1a:



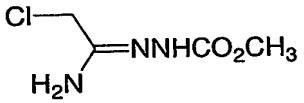
1a

is disclosed wherein the hydrochloride salt of a compound of formula 2a:



2a

is reacted in the presence of an inorganic base and toluene with a compound of the formula 3:



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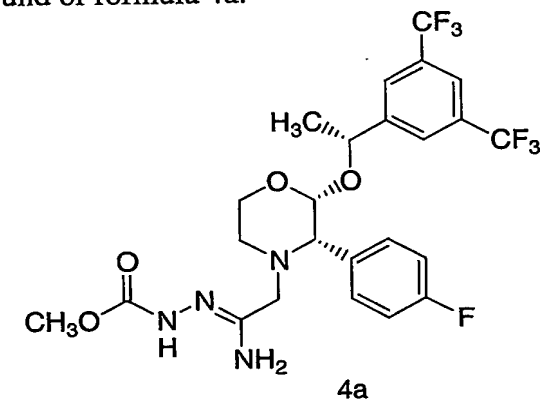
to produce the compound of formula 4a:



4a ;

(b) washing with an aqueous phase and

(c) cyclizing at a temperature of 140-150°C to produce the compound of formula 1a.

The washing step described herein typically uses an aqueous phase, e.g., water, and may optionally contain a salt. Representative examples of salts that are useful herein include KCl, $KHCO_3$, K_2CO_3, Na_2CO_3, $NaHCO_3$, NaCl and similar such salts. KCl is the preferred salt.

In another aspect of the invention, the process is further comprised of a drying step prior to cyclization.

As used herein the term "inorganic base" refers to compounds such as sodium carbonate, cesium carbonate, sodium hydroxide, potassium hydroxide, potassium carbonate and the like. More particularly, the preferred inorganic base is potassium carbonate.

More particularly, the present invention relates to the process described above wherein compound 2 or 2a is reacted with compound 3 in the presence of an inorganic base, toluene and a polar aprotic solvent. As used herein, the term "polar aprotic solvent" refers to a solvent that neither donates or accepts protons, and is, for example, selected from the group consisting of: dimethylformamide (DMF), dimethylsulfoxide (DMSO), N-methylpyrrolidone (NMP), acetonitrile (MeCN), N,N-dimethylacetamide (DMAC) and hexamethylphosphoramide (HMPA).

The process described herein is surprisingly efficient, minimizing the production of a mixture of isomers, and thus increasing productivity and purity. The subject process also minimizes the use of toxic solvents.

The 2-[1-[3,5-bis(trifluoromethyl)phenyl]ethoxy-3-(4-fluorophenyl)-1,4-oxazine starting material 2 and (2R, 2-alpha-R, 3a)-2-[1-[3,5-bis(trifluoromethyl)-phenyl]ethoxy-3-(4-fluorophenyl)-1,4-oxazine starting material 2a may be obtained in accordance with PCT WO 01/94324 A1 (published December 13, 2001) and US 2002/0052494 A1 (published May 2, 2002), or using modifications thereof. The starting material may be used directly or following purification. Purification procedures include crystallization, distillation, normal phase or reverse phase chromatography. The following example is provided for purposes of illustration and is not intended to limit the disclosed invention.

EXAMPLE 1

[2*R*-[2α(*R**),3α]]-5-[[2-[1-[3,5-bis(trifluoromethyl)phenyl]ethoxy]-3-(4-fluorophenyl)-4-morpholinyl]methyl]-1,2-dihydro-3*H*-1,2,4-triazol-3-one

A mixture of the starting material as the hydrochloride salt of (2R, 2-alpha-R, 3a)-2-[1-[3,5-bis(trifluoromethyl)phenyl]ethoxy-3-(4-fluorophenyl)-1,4-oxazine (2a) (1.00 kg; 2.11 mol) and potassium carbonate (1.02 kg; 7.39 mol) in DMSO (2.2 L) and toluene (1.0 L) was cooled to 15°C. A slurry of amidrazone 3 (367 g; 2.22 mol) in toluene (1.5 L) was added. The mixture was stirred and then partitioned between toluene (4.0 L) and water (5.0 L). The phases were separated at 40°C. The organic layer (containing 4a) was washed with water (5.0 L) at 40°C and

then partially concentrated at atmospheric pressure, providing intermediate 4a, which is used in the next step without isolation. The resulting solution containing intermediate 4a was heated to 140°C for 3 h and then allowed to cool to RT. The solids were filtered and dried *in vacuo* at 40 °C. The product (1.00 kg) was dissolved in methanol (10.0 L) and 50 g of Darco was added. The mixture was heated at 60°C for 1 h and then filtered at this temperature. The filtrates were allowed to cool slowly to RT. Water (5.0 L) was added slowly over 1 h. The slurry was cooled to 5 °C and the solids were filtered and dried *in vacuo* at 40 °C to yield 0.96 kg (85% overall yield) of the product [2*R*-[2α(*R**),3α]]-5-[[2-[1-[3,5-bis(trifluoromethyl)phenyl]-ethoxy]-3-(4-fluorophenyl)-4-morpholinyl]methyl]-1,2-dihydro-3*H*-1,2,4-triazol-3-one (i.e. 5-[[2(R)-[1(R)-[3,5-bis(trifluoromethyl)phenyl]ethoxy]-3(S)-(4-fluorophenyl) -4-morpholinyl]methyl]-1,2-dihydro-3H-1,2,4-triazol-3-one).

Intermediate 4a: $[\alpha]_D^{25}$ = +84° (c=1.02, methanol); ^{1}H NMR (400 MHz, CDCl3) δ 7.64 (s, 2H), 7.34 (br t, *J* ~ 7, 2H), 7.16 (s, 1H), 7.03 (t, *J* = 8.4, 2H), 5.8 (very br s, 2H), 4.88 (q, *J* = 6.6, 1H), 4.33 (d, *J* = 2.8, 1H), 4.24 (td, *J* = 11.6, 2.0, 1H), 3.77 (s, 2H), 3.66 (ddd, *J* = 11.6, 3.2, 1.6, 1H), 3.46 (d, *J* = 2.8, 1H), 3.31 (d, *J* = 14.5, 1H), 2.96 (br d, *J* = 11.6, 1H), 2.59 (d, *J* = 14.5, 1H), 2.50 (td, *J* = 12.1, 3.2, 1H), 1.47 (d, *J* = 6.6, 3H). Anal. Calc. for $C_{24}H_{25}F_7N_4O_4$: C, 50.89; H, 4.45; F, 23.48; N, 9.89. Found: C, 50.48; H, 4.40; F, 23.43; N, 9.84. Final product 1a: Mp: 255 °C; $[\alpha]_D^{25}$ = +69° (c=1.00, methanol); ^{1}H NMR (400 MHz, CD_3OD) δ 7.70 (s, 1H), 7.51 (m, 2H), 7.32 (s, 2H), 7.04 (t, *J* = 8.7, 2H), 4.94 (q, *J* = 6.3, 1H), 4.35 (d, *J* = 2.8, 1H), 4.28 (td, *J* = 11.5, 2.8, 1H), 3.66 (ddd, *J* = 11.5, 3.3, 1.6, 1H), 3.54 (d, *J* = 14.3, 1H), 3.48 (d, *J* = 2.8, 1H), 2.88 (br d, *J* = 11.9, 1H), 2.86 (d, *J* = 14.3, 1H), 2.49 (td, *J* = 11.9, 3.6, 1H), 1.44 (d, *J* = 6.3, 3H); ^{13}C NMR (100 MHz, CD_3OD) δ 164.1 (d, *J* = 245.9), 158.7, 147.6, 147.0, 134.1 (d, *J* = 3.1), 132.7 (d, *J* = 33.4), 132.4 (d, *J* = 8.0), 127.8 (m), 124.6 (q, *J* = 272.0), 122.3 (m), 116.1 (d, *J* = 21.6), 97.1, 73.7, 70.5, 60.4, 53.6, 52.2, 24.7. Anal. Calc. for $C_{23}H_{21}F_7N_4O_3$: C, 51.69; H, 3.96; F, 24.88; N, 10.48. Found: C, 51.50; H, 3.82; F, 24.73; N, 10.44. HRMS: 534.1480 (meas.); 534.1502 (calc. for $C_{23}H_{21}F_7N_4O_3$).

All patents and patent publications cited herein are incorporated by reference in their entirety. While the invention has been described and illustrated with reference to certain particular embodiments thereof, those skilled in the art will appreciate that various adaptations may be made without departing from the spirit and scope of the invention.